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JUL 80 R ABBUNDI, A E CLARK

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MAGNETOSTRICTION OF LAVES PHASE RARE EARTH-Ni₂ COMPOUNDS

BY R. ABBUNDI, A. E. CLARK

RESEARCH AND TECHNOLOGY DEPARTMENT

1 JULY 1980

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20. ABSTRACT (Continue on reverse side if necessary and identify by block number) The magnetostriiction of the RNi ₂ compounds (R = Tb, Dy, Ho, Er and Tm) was measured as a function of applied field from T = 5 K to above the Curie temperatures. A direct comparison is made between these results and the huge magnetostriiction observed at cryogenic temperatures in the RFe ₂ and RCo ₂ compounds. The largest magnetostriiction in the RNi ₂ series occurs in TbNi ₂ . At T = 5 K the magnetostriiction is fairly well saturated and results in a value of $\sigma_{11} = 2270 \times 10^{-6}$ at H = 25 kOe, while extrapolation to infinite field yields $\sigma_{11} = 2340 \times 10^{-6}$. DyNi ₂ also displays a tendency to saturate at the higher fields.		

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→ at lower temperatures with $\lambda_{11} - \lambda_{12} = -765 \times 10^{-6}$ at 25 kOe and -840 ppm at $H = 6$. The compounds HoNi_2 , ErNi_2 , and TmNi_2 all exhibit large strains at $T = 5$ K although fail to saturate with available fields. At 25 kOe the magnetostriction for these compounds was found to be 586×10^{-6} , 415×10^{-6} and -552×10^{-6} respectively. ←

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FOREWORD

The magnetostriction study reported here is part of a research program undertaken to determine the nature of the magnetostriction in the rare earth-intermetallic compounds. In this paper is detailed the temperature dependence of the magnetostriction for a series of rare earth-Ni₂ compounds. Studies were made as a function of applied field from T = 5 K to temperatures above the Curie points.

The largest magnetostriction in the RNi₂ series occurs in TbNi₂. At T = 5 K the magnetostriction is fairly well saturated and results in a value of $\lambda_{||} - \lambda_{\perp} = 2270 \times 10^{-6}$ at H = 25 kOe, while extrapolation to infinite field yields 2540×10^{-6} . DyNi₂ also displays a tendency to saturate at the higher fields at lower temperature with $\lambda_{||} - \lambda_{\perp} = -765 \times 10^{-6}$ at 25 kOe and -840 ppm at H = ∞ . The compounds HoNi₂, ErNi₂, and TmNi₂ all exhibit large strains at T = 5 K although fail to saturate with available fields. At 25 kOe the magnetostriction for these compounds was found to be -386×10^{-6} , -415×10^{-6} and -552×10^{-6} respectively.

The study was carried out in the Solid State Branch of the Radiation Division as part of the research program on magnetostrictive material. The research was sponsored by the Office of Naval Research (PO-4-0081, NR 039-110) and the NSWC Independent Research Program (IR-011).

B. F. DESAVAGE
By direction

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INTRODUCTION

Intermetallic compounds of the rare earths with the 3d transition metals of Fe, Co and Ni have been the subject of extensive investigations at many laboratories. Many interesting effects have been observed in ultrasonic and magnetostriction measurements in the cubic Laves phase RTM_2 compounds, where large changes in the elastic moduli as well as huge magnetostriction constants occur.¹⁻⁷ In this work we report on the magnetostriction of the RNi_2 compounds ($R = Tb, Dy, Ho, Er$ and Tm).

Bulk magnetization measurements⁸ on the RNi_2 series have shown that the Curie temperatures are in the cryogenic region with $T_c < 90$ K. Both YNi_2 and $LuNi_2$ were found to exhibit only a Pauli paramagnetic behavior down to 4.2 K. Thus it appears that Ni possesses no moment of its own in these compounds. This behavior is unlike that observed in the RFe_2 and RCo_2 compounds. In the case of the RFe_2 series the 3d transition element when combined with the heavy lanthanide elements Tb through Tm possesses a moment of approximately $1.6 \mu_B$ /Fe atom, with only a relatively small variation dependent upon the rare earth.^{9,10} In the RCo_2 compounds the Co moment is approximately $1 \mu_B$ /Co atom.¹¹ The magnetization results⁸ further showed that $GdNi_2$ at $T = 0$ possesses a saturation moment of $7.1 \mu_B$, very nearly equal to the gJ value of the rare earth. However, the moments of the remaining RNi_2 compounds are substantially reduced from their theoretical gJ values. This discrepancy has been attributed to crystal field effects on the rare earth which partially quench the orbital angular momentum.¹²

Mössbauer effect measurements¹³ on the RNi_2 compounds have shown that the easy axis of magnetization follows the same sequence as that observed for both the RFe_2 ¹⁴ and RCo_2 ¹⁵ series. $DyNi_2$ and $HoNi_2$ magnetize along a [100] direction, while $TbNi_2$ and $ErNi_2$ along a [111] direction. Although no results were reported on $TmNi_2$, it is assumed that $TmNi_2$ has its easy axis of magnetization along a [111] direction, similar to $TmFe_2$.¹⁶

The magnetostriction measurements were performed as a function of field from $T = 5$ K to above the Curie temperatures. These results can be directly compared with the huge magnetostriction which has been found at cryogenic temperatures in both the RFe_2^{3-5} and $RCo_2^{6,7}$ series.

EXPERIMENTAL RESULTS

The highly anisotropic 4f electron charge cloud of the Tb^{3+} ion produces enormous magneto-strains in TbFe_2 ³ and TbCo_2 .^{6,7} The high Curie temperature of TbFe_2 ($T_c = 710$ K) allows this huge magnetostriction of $\lambda_{111}(0 \text{ K}) = 4400 \times 10^{-6}$ at $T = 0$ to remain large at room temperature, where $\lambda_{111}(300 \text{ K}) = 2500 \times 10^{-6}$. This strain is the largest room temperature magnetostriction of any known material. In TbCo_2 a similarly huge rhombohedral distortion develops below $T_c = 240$ K, also reaching 4400×10^{-6} at 4.2 K.

Gignoux et al.⁷ state that TbNi_2 , however, behaves quite differently. Rhombohedral distortions (λ_{111}), measured by x-ray diffraction, were found to be ≈ 0 . They state that this small value is consistent with a smaller molecular field than found in either TbFe_2 or TbCo_2 resulting in a smaller value of $\langle 0_2^0 \rangle$. Our results on the magnetostriction of TbNi_2 do not confirm this low measured value. Instead, we find that the strain in TbNi_2 is about 50% of that in TbFe_2 and TbCo_2 . Figure 1 shows the magnetostriction at $T = 5$ K as a function of applied field for TbNi_2 and DyNi_2 . The temperature dependence of the magnetostriction at $H = 25$ kOe for TbNi_2 and DyNi_2 is shown in Figure 2. At $T = 5$ K the magnetostriction of TbNi_2 is fairly well saturated and results in a value of $\lambda_{||} - \lambda_{\perp} = 2270 \times 10^{-6}$ at $H = 25$ kOe, while extrapolation to infinite field yields 2340×10^{-6} .

DyNi_2 also displays a tendency to saturate with $\lambda_{||} - \lambda_{\perp} = -765 \times 10^{-6}$ at 25 kOe. The magnitude of this strain while comparable to that observed in DyCo_2 ⁶ implies a substantially larger $|\lambda_{100}|$ than was found in DyFe_2 , where $|\lambda_{100}|$ remained small through the entire temperature range.³

The field dependence of the magnetostriction at $T = 5$ K for HoNi_2 , ErNi_2 and TmNi_2 is shown in Figure 3. A peak in the magnetostriction of HoNi_2 was observed near $H = 5$ kOe. We attribute this behavior to the large magnetocrystalline anisotropy which characterizes the RTM_2 compounds.^{7,9,17-20} In the case of HoNi_2 , as the field is increased above 5 kOe, the moments are tilted away from their highly magnetostrictive easy [100] axes, decreasing the net strain. This behavior is only

observed at the lower temperatures. For $T > 11$ K the rapid decrease in anisotropy with increasing temperature results in the usual behavior of increasing $|\lambda|$ with increasing field. The magnitude of the peak strain in HoNi_2 at $T = 5$ K of -533 ppm is comparable to λ_{100} measured in HoFe_2 where $\lambda_{100}(0 \text{ K}) = -745$ ppm.⁵ The temperature dependence of the magnetostriction at $H = 25$ kOe for HoNi_2 is shown in Figure 4.

The large anisotropy in these materials is again reflected in the failure of ErNi_2 to reach saturation at 5 K, as shown in Figure 3. However, the magnetostriction is still large with $\lambda_{||} - \lambda_{\perp} = -415 \times 10^{-6}$ at $H = 25$ kOe. Figure 5 shows the temperature dependence of the strain at 25 kOe, from 5 K to above the Curie temperature.

A comparison of the field dependence of the magnetostriction at 5 K between TmNi_2 (see Figure 3) and the other compounds shows a substantially different behavior. The magnetostriction is far from saturation. Farrell and Wallace, in their bulk magnetization measurements,⁸ report that TmNi_2 may not be ferromagnetic at $T = 2$ K. They could not determine whether TmNi_2 was displaying VanVleck paramagnetism or the onset of ferromagnetism. However, in view of those findings, it appears that, at least at 5 K, TmNi_2 is not ferromagnetic and thus that accounts for the absence of any spontaneous magnetostriction at this temperature. Nevertheless at $H = 25$ kOe the strain is becoming quite large with $\lambda_{||} - \lambda_{\perp} = -552 \times 10^{-6}$. In theory the saturation magnetostriction of TmNi_2 should be of comparable magnitude to that found in TbNi_2 . The temperature dependence of the magnetostriction for TmNi_2 at 25 kOe is shown in Figure 6.

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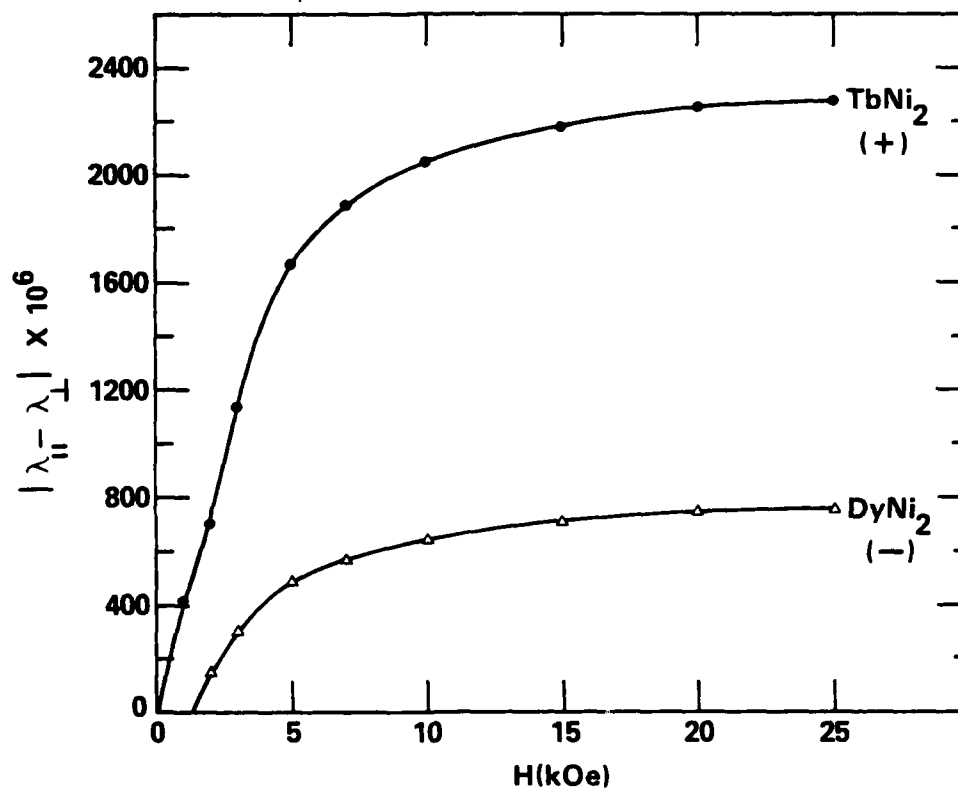


FIGURE 1 MAGNETOSTRICTION AT T=5K AS A FUNCTION OF APPLIED FIELD FOR TbNi₂ AND DyNi₂. THE MAGNETOSTRICTION OF TbNi₂ IS POSITIVE (+), WHILE DyNi₂ IS NEGATIVE (-).

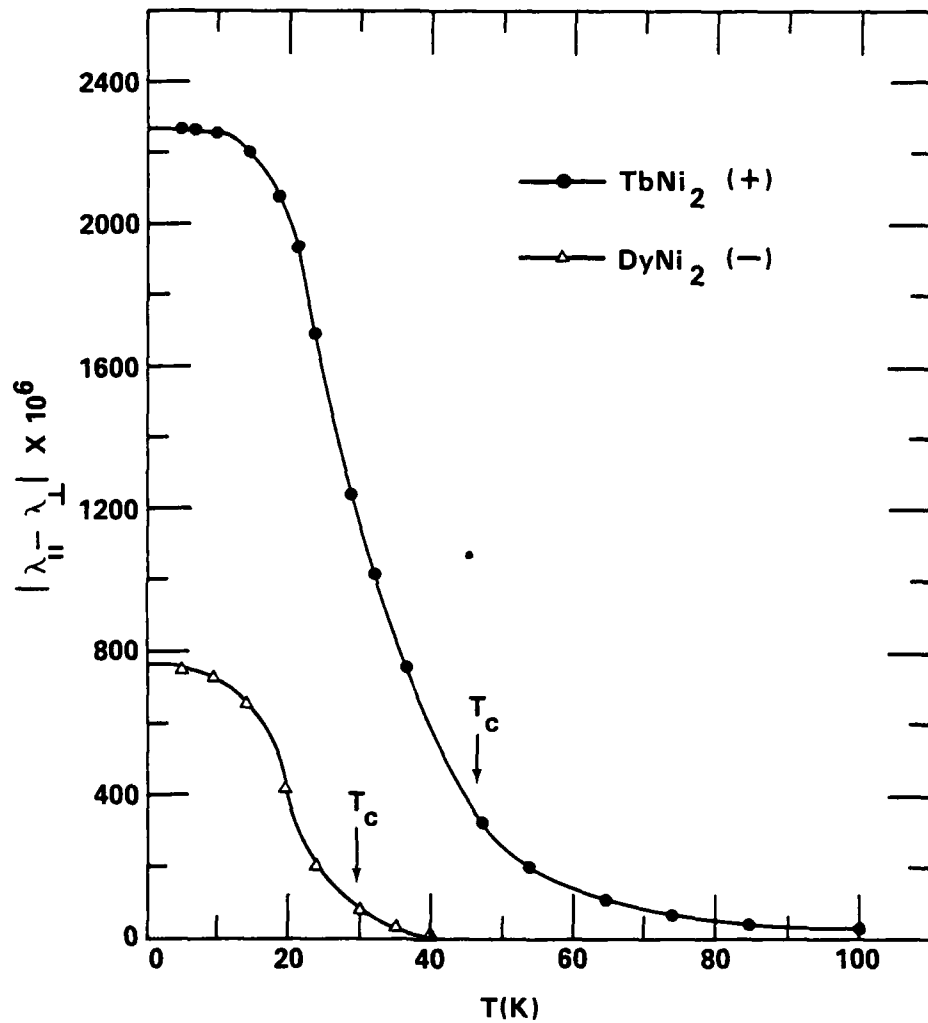


FIGURE 2 TEMPERATURE DEPENDENCE OF THE MAGNETOSTRICTION AT $H=25\text{kOe}$ FOR TbNi_2 AND DyNi_2 . THE CURIE TEMPERATURES WERE TAKEN FROM BULK MAGNETIZATION MEASUREMENTS (REF. 8).

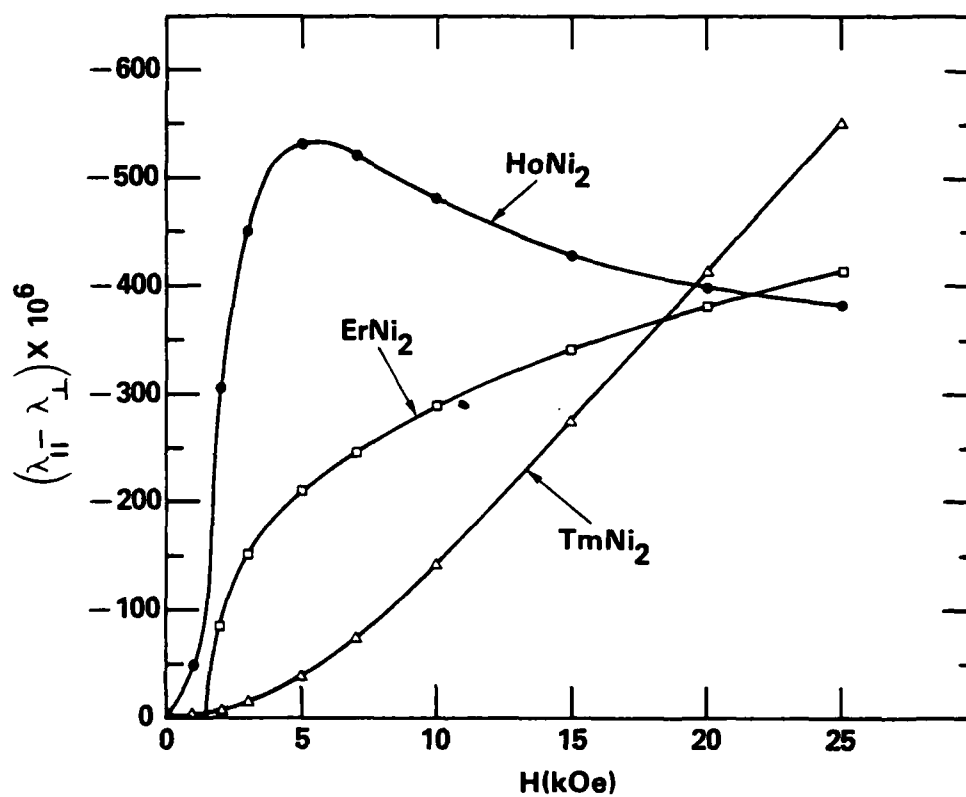


FIGURE 3 MAGNETOSTRICTION AT T=5K AS A FUNCTION OF APPLIED FIELD FOR HoNi_2 , ErNi_2 AND TmNi_2 .

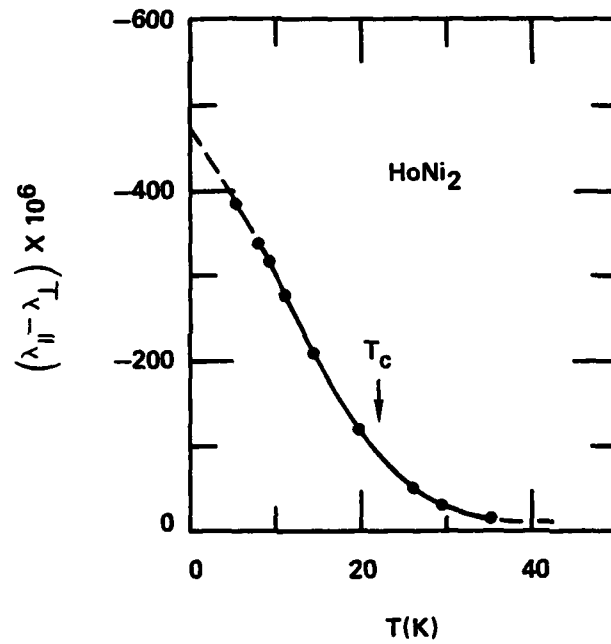


FIGURE 4 TEMPERATURE DEPENDENCE OF THE MAGNETOSTRICTION AT $H=25\text{kOe}$ FOR HoNi_2 . THE CURIE TEMPERATURE WAS TAKEN FROM BULK MAGNETIZATION MEASUREMENTS (REF. 8).

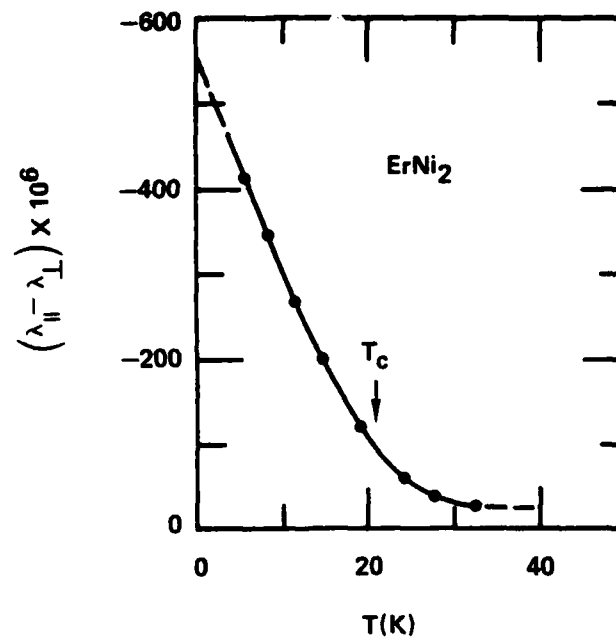


FIGURE 5 TEMPERATURE DEPENDENCE OF THE MAGNETOSTRICTION AT $H=25\text{kOe}$ FOR ErNi_2 . THE CURIE TEMPERATURE WAS TAKEN FROM BULK MAGNETIZATION MEASUREMENTS (REF. 8).

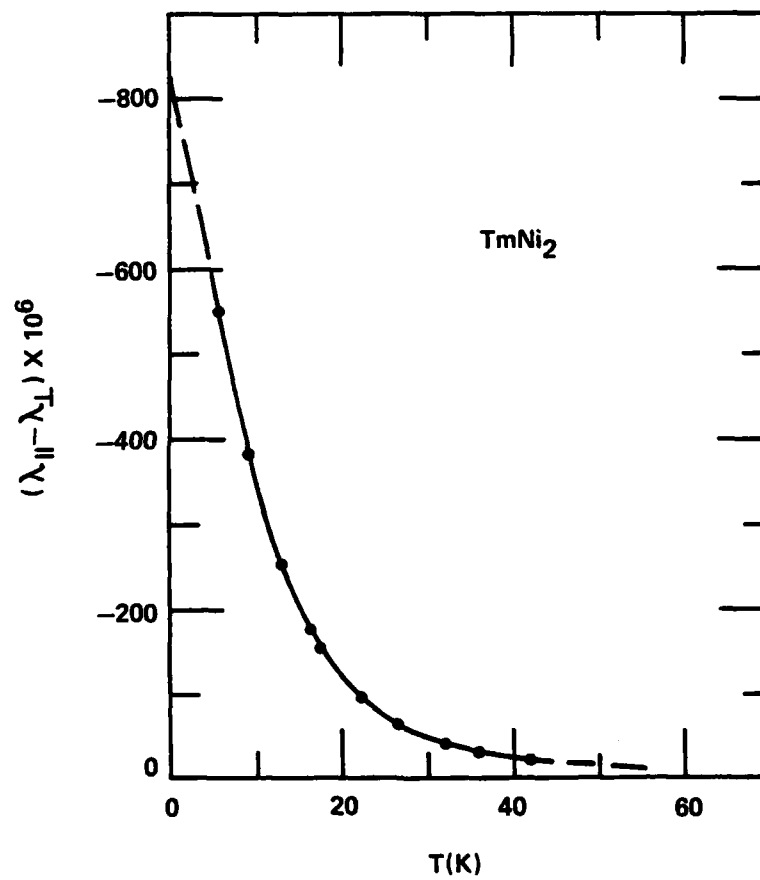


FIGURE 6 TEMPERATURE DEPENDENCE OF THE MAGNETOSTRICTION
AT $H=25\text{kOe}$ FOR TmNi_2

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